DATE: March 25,1998

SUBJECT: Results for Water Supply Performance

Evaluation Study 40 (WS040)

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National Water Quality Assurance Programs Branch

Ecological Exposure Research Division

TO: Designated USEPA and State WS Study Coordinators,

and Selected Individual Laboratory Addressees

THRU: Raymond J. Wesselman, Chief

National Water Quality Assurance Programs Branch

Ecological Exposure Research Division

I. DISCUSSION OF WS040 CHEMISTRY RESULTS

WS040 has been completed by the National Exposure Research Laboratory - Cincinnati (NERL-Cincinnati) as the first FY98 study for evaluation of U.S. Environmental Protection Agency (USEPA) regional laboratories, the state laboratories and other selected laboratories involved in chemical analyses covered under the Drinking Water Laboratory Certification Program. As seen in the enclosed summaries, the majority of laboratories produced acceptable results from these samples; 88.6 percent of the usable results reported were acceptable. The results relating to your interests are also enclosed.

No analyte had a failure rate over 30 percent and two analytes were not evaluated:

1) The two low-level turbidity analytes were NOT EVALUATED for this study. These data were collected for information purposes and were never intended to be evaluated as a part of WS040. A separate memorandum summarizing the low-level turbidity data has been enclosed for your information and possible distribution.

Within the data reported for VOC #2, the only qualitative-challenge group in this study, the missing analyte with the highest false positive rates was 4-chlorotoluene at about 4 percent of those reporting data for VOC #2.

This memorandum highlights the analytes that had a high rate of "NOT ACCEPT." analytical responses. It is each laboratory management's responsibility to investigate their "NOT ACCEPT." results to discover their own specific problems.

For those interested in the aroclor present in the PCB sample, it was 1242.

II. GENERAL INFORMATION

In response to 40 CFR Part 141 modifications, the following acceptance limits were used at all concentration levels unless otherwise specified:

<u>Analyte</u>	Acceptance Limits
Antimony Barium Beryllium Cadmium Chromium Copper Lead Mercury Nickel Selenium Thallium	True Value (TV) + 30%, for TV > 6 μg/L TV + 15%, for TV > 150 μg/L TV + 20%, for TV > 1 μg/L TV + 15%, for TV > 2 μg/L TV + 15%, for TV > 10 μg/L TV + 10%, for TV > 50 μg/L TV + 30%, for TV > 5 μg/L TV + 30%, for TV > 10 μg/L TV + 20%, for TV > 2 μg/L TV + 30%, for TV > 2 μg/L TV + 30%, for TV > 2 μg/L TV + 20%, for TV > 10 μg/L TV + 20%, for TV > 10 μg/L TV + 30%, for TV > 10 μg/L TV + 30%, for TV > 2 μg/L
<u>Analyte</u>	Acceptance Limits
Nitrate Nitrite Fluoride Total Cyanide Alachlor Atrazine Chlordane	TV $\frac{+}{+}$ 10%, for TV $\frac{>}{-}$ 0.4 mg/L TV $\frac{+}{+}$ 15%, for TV $\frac{>}{-}$ 0.4 mg/L TV $\frac{+}{+}$ 10%, for TV between 1 and 10 mg/L TV $\frac{+}{+}$ 25%, for TV $\frac{>}{-}$ 0.1 mg/L TV $\frac{+}{+}$ 45% TV $\frac{+}{+}$ 45%

<u>Analyte</u>	Acceptance Limits		
Endrin Heptachlor Heptachlor epoxide Lindane Methoxychlor Toxaphene Carbofuran	TV + 30% TV + 45%		
<u>Analyte</u>	Acceptance Limits		
2,4-D 2,4,5-TP (Silvex) Pentachlorophenol Decachlorobiphenyl THMs DBCP EDB Vinyl chloride all other regulated	True Value (TV) + 50% TV + 50% TV + 50% TV + 100% TV + 20% TV + 40% TV + 20%, for TV <10 µg/L TV + 20%, for TV > 10 µg/L		
(Benzene, Carbon Tetrachloride, Chlorobenzene, 1,2-Dichlorobenzene, 1,4-Dichlorobenzene, 1,2-Dichloroethane, 1,1-Dichloroethylene, Cis-1,2-Dichloroethylene, Trans-1,2-Dichloroethylene, Dichloromethane, 1,2-Dichloropropane, Ethylbenzene, Styrene, Tetrachloroethylene, Toluene, 1,1,1-Trichloroethane, 1,1,2-Trichloroethane, Trichloroethylene,			

For all other analytes and/or concentrations: A statistical 95% prediction interval was used based on the statistics of analytical results from USEPA and state laboratories.

1,2,4-Trichlorobenzene, and Total Xylenes)

The report for each participating Office of Research and Development (ORD) laboratory is sent to the Laboratory Director. Reports for participating contract/grant laboratories are sent to the responsible QA Officer. Regional and state coordinators will find enclosed a personal computer disk containing the study files of interest to them, one copy of the report for each of their participating laboratories and their part of the study participant list. The addressees are responsible for any

additional distribution of study results that may be necessary to properly inform state agencies and other participants.

Regarding the procedure that we have established for formal correction of data entry errors, laboratories are responsible for reporting any data entry errors in their report. These errors must to be reported as soon as possible, however, we will accept errors reported up to four (4) months from the date on the cover memorandum used by us for the distribution of individual laboratory reports at the conclusion of the study. If confirmed in our records, errors received before the four-month deadline will be corrected in our study file and report pages will be corrected and reissued to the laboratory and the coordinators that nominated that laboratory. After the four months, PC disks containing the corrected study results will be distributed to each coordinator.

For each laboratory, the <u>Participant List</u> shows all the coordinators that requested that laboratory's participation and identifies the coordinator with primary responsibility for informing the laboratory. If your region/state is the only requesting office, or if your region/state is listed after "samples thru" in the <u>Participant List</u>, it is your responsibility to provide that laboratory with a copy of their evaluation report. Every coordinator is responsible for seeing that any laboratory they requested receives any study summaries, true values, acceptance limits, etc., that the laboratory may request. Requestors for such study information reaching NERL-Cincinnati or ManTech, will be instructed to contact their study coordinator(s) for this information.

In addition, each Regional Coordinator is responsible for assuring that each of their states receives all appropriate study information.

For each "NOT ACCEPT." performance evaluation received, the laboratory should determine the cause(s) and make the procedural changes necessary to improve future data quality.

Thank you for your continued cooperation in these studies. If you have any questions about or problems with the study or the reports, please do not hesitate to contact me at (513) 569-7216.